

3-10-82

BB-1143  
TXR-2807

UNITED STATES ENVIRONMENTAL PROTECTION AGENCY

DATE  
SUBJECT: EPA File Symbol 476-EERE  
LC-6641 Selective Herbicide

002807

FROM: Deloris F. Graham *DFG 3/10/82*  
PHB/TSS  
TO: Robert J. Taylor  
Product Manager (25)

Applicant: Stauffer Chemical Company  
1200 South 47th Street  
Richmond, California 94804

Active Ingredient:  
Sutan: S-ethyl diisobutylthiocarbamate . . . . . 53%  
Atrazine: 2-chloro-4-(alkylamino)-6-(isopropylamino)-s-triazine . . 12.98%  
Related Triazines . . . . . 0.27%  
Inert Ingredients . . . . . 33.75%

Background: Submitted Acute Oral, Acute Dermal, Eye Irritation and Skin Irritation Studies. Studies conducted by Stauffer Chemical Company. Data under Accession Number 246784. Combined cite-all and alternate method of support.

Recommendations:

- (1) PHB/TSS finds these data acceptable to support conditional registration of this product. However, for future submission please note:
  - a. In the Eye Irritation Study, individual scoring for corneal opacity, iris irritation and conjunctive irritation per animal must be submitted.
  - b. In the Primary Dermal Irritation Study, 4 sites (2 abraded and 2 intact) per animal must be used.
- (2) The appropriate signal word is DANGER.

Label:

- (1) The appropriate signal word is DANGER and must appear on center front panel of label.
- (2) The following statement must precede the "Harmful if swallowed" statement:

"Corrosive, cause irreversible eye damage".

*1 Jlt*

- (3) The statement "Keep out of lakes, streams, and ponds" must be revised to read "Do not apply directly to water."

Review:

- (1) Acute Oral Toxicity Study: Stauffer Chemical Company; Report #T-10880; February 4, 1982.

Procedure: 7 groups, 5 of which consisted of 10M and 10F, and 2 of which consisted of 20M and 10F, each received one of the following doses: 2000, 2200, 2500, 2800, 3200, 4000 and 5000 mg/kg. Observations made daily for 14 to 15 days after treatment. Necropsy performed on all animals.

Results: at 2200 mg/kg, 1/10M died; at 2500 mg/kg, 12/20M and 2/10F died; at 2800 mg/kg, 8/20M died; at 3200 mg/kg, 6/10M and 2/10F died; at 4000 mg/kg, 9/10M and 4/10F died; at 5000 mg/kg, 10/10M and 9/10F died.

Toxic signs included moderate depression, ptosis, chromodacryorrhea, severe depression, red facial stains, yellow discoloration around the anogenital region, salivation, reddened lungs, white solid mass in the urinary bladder and dehydration.

Necropsy revealed white fluid in the stomach or gastrointestinal tract; brown or red-brown foci on the cardia portion of the stomach; stomach wall appeared thin; gray-tan discoloration of the liver; similar discoloration of the spleen, left kidney and left adrenal; reddened lungs and gastrointestinal tract; reddened jejunum and scotal skin; red to yellow discoloration around the muzzle.

LD<sub>50</sub> for males was 2780 mg/kg with 95% confidence limits between 2490 and 3104 mg/kg. LD<sub>50</sub> for females was 3600 mg/kg with 95% confidence limits between 3167 and 4093 mg/kg.

Study Classification: Core Guideline Data

Toxicity Category: III-CAUTION

- (2) Acute Dermal Toxicity Study: Stauffer Chemical Company; Report #T-10675; January 29, 1982.

Procedure: 4M and 4F rabbits received 2000 mg/kg of the test material under occlusive wrap for 24 hour exposure. Half the animals had abraded skin. Observations made for 14 days after treatment. Necropsy performed on all animals.

Results: No mortalities or toxic signs. No abnormalities at necropsy. Mild to moderate erythema and edema noted. LD<sub>50</sub> greater than 2000 mg/kg.

Study Classification: Core Guideline Data

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Toxicity Category: III - CAUTION

- (3) Eye Irritation Study: Stauffer Chemical Company; Report #T-10675;  
January 29, 1982.

Procedure: 9 rabbits received 100 mg of the test material in one eye each. The eyes of three of the rabbits were washed 20-30 seconds posttreatment. Observations made at 24, 48, 72 hours and at 4 and 7 days. If irritation was present at 7 days an additional observation made every 3-4 days until injury subsided or was found to be irreversible.

Results: 6/6 of the unwashed group showed severe corneal opacity and moderate to severe redness, chemosis, discharge. 4/6 moderate to severe iris irritation. At day 7, minimal remission of these symptoms. At 14 days, 1/6 moderate irritation. Pannus and corneal ulcer observed through day 28; clear by day 31.

2/3 of the washed groups showed mild opacity and 1/3 of the washed group moderate to severe opacity. 1/3 showed moderate irritation of the iris. 3/3 showed moderate to severe redness and chemosis and mild to moderate discharge. Complete reversal of these symptoms occurred by seven days.

Study Classification: Core Minimum Data. Individual scoring for corneal opacity, iris irritation and conjunctive irritation per animal must be submitted.

Toxicity Category: I - DANGER

- (4) Primary Dermal Irritation Study: Stauffer Chemical Company; Report #T-10675; January 29, 1982

Procedure: 6 rabbits received 0.5g of the test material at one intact and one abraded skin site per animal under occlusive wrap for 24 hour exposure. Observations were made at 24 and 72 hours.

Results: At 24 hours, 6/6 animals had mild to moderate erythema (4/6=1, 2/6=2) and 4/6 edema (3/6=1, 1/6=2). At 72 hours, 2/6 showed mild to moderate erythema (1/6=1, 1/6=2) and 1/6 mild edema (1/6=1). Primary irritation score was 1.3.

Study Classification: Core Minimum Data. 4 sites, 2 abraded and 2 intact, per animal must be used.

Toxicity Category: III - CAUTION

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  - ☐ Identity of product impurities
  - ☐ Description of the product manufacturing process
  - ☐ Description of product quality control procedures
  - ☐ Identity of the source of product ingredients
  - ☐ Sales or other commercial/financial information
  - ☒ A draft product label
  - ☐ The product confidential statement of formula
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